AMENDMENTS TO THE CLAIMS

- 1. (Previously presented) A mixture, comprising
 - a) a compound of the formula I

in which

- X is halogen, C₁-C₄-alkyl or trifluoromethyl;
- m is 0 or 1;
- Q is C(=CH-CH₃)-COOCH₃, C(=CH-OCH₃)-COOCH₃, C(=N-OCH₃)-COOCH₃ or N(-OCH₃)-COOCH₃;
- A is -O-B, -CH₂O-B, -OCH₂-B, -CH=CH-B, -C \equiv C-B, -CH₂O-N=C(R¹)-B or -CH₂O-N=C(R¹)-C(R²)=N-OR³, where

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B is phenyl, naphthyl, 5-membered or 6-membered hetaryl or 5-membered or 6-membered heterocyclyl which contains one to three nitrogen atoms and/or one oxygen or sulfur atom or one or two oxygen and/or sulfur atoms, where the ring systems are unsubstituted or substituted by one to three radicals R^a:

R^a is cyano, nitro, amino, aminocarbonyl, aminothiocarbonyl, halogen, C₁-C₆-alkyl, C₁-C₆-haloalkyl, C₁-C₆-alkylcarbonyl, C₁-C₆-alkylsulfonyl, C₁-C₆-alkylsulfoxyl, C₃-C₆-cycloalkyl, C₁-C₆-alkoxy, C₁-C₆-alkyloxycarbonyl, C₁-C₆-alkylthio, C₁-C₆-alkylamino, di-C₁-C₆-alkylamino, C₁-C₆-alkylaminocarbonyl, di-C₁-C₆-alkylaminocarbonyl, di-C₁-C₆-alkylaminothiocarbonyl, di-C₁-C₆-alkylaminothiocarbonyl,

C₂-C₆-alkenyl, C₂-C₆-alkenyloxy, phenyl, phenoxy, benzyl, benzyloxy, 5- or 6-membered heterocyclyl, 5- or 6-membered hetaryl, 5- or 6-membered hetaryloxy, C(=NOR')-OR" or OC(R')₂-C(R")=NOR",

where the cyclic radicals for their part are unsubstituted or substituted by one to three radicals R^b:

R^b is cyano, nitro, halogen, amino, aminocarbonyl, aminothiocarbonyl, C₁-C₆-alkyl, C₁-C₆-haloalkyl, C₁-C₆-

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alkylsulfonyl, C₁-C₆-alkylsulfoxyl, C₃-C₆-cycloalkyl, C₁-C₆-alkoxy, C₁-C₆-alkoxy, C₁-C₆-alkoxy, C₁-C₆-alkylamino, C₁-C₆-alkylamino, C₁-C₆-alkylamino, C₁-C₆-alkylaminocarbonyl, di-C₁-C₆-alkylaminocarbonyl, C₁-C₆-alkylaminothiocarbonyl, di-C₁-C₆-alkylaminothiocarbonyl, C₂-C₆-alkylaminothiocarbonyl, C₂-C₆-alkenyloxy, C₃-C₆-cycloalkyl, C₃-C₆-cycloalkyl, C₃-C₆-cycloalkyl, phenyl, phenoxy, phenylthio, benzyl, benzyloxy, 5- or 6-membered heterocyclyl, 5- or 6-membered hetaryl, 5- or 6-membered hetaryloxy or C(=NOR')-OR";

- R' is hydrogen, cyano, C_1 - C_6 -alkyl, C_3 - C_6 -cycloalkyl or C_1 - C_4 -haloalkyl;
- R" is hydrogen, C_1 - C_6 -alkyl, C_3 - C_6 -alkenyl, C_3 - C_6 -alkinyl, C_1 - C_4 -haloalkyl, C_3 - C_6 -haloalkenyl or C_3 - C_6 -haloalkinyl;
- R¹ is hydrogen, cyano, C₁-C₄-alkyl, C₁-C₄-haloalkyl, C₃-C₆-cycloalkyl, C₁-C₄-alkoxy;
- R² is phenyl, phenylcarbonyl, phenylsulfonyl, 5- or 6-membered hetaryl, 5- or 6-membered hetarylcarbonyl or 5- or 6-membered hetarylsulfonyl,

where the ring systems are unsubstituted or substituted by one to three radicals R^a,

is C_1 - C_{10} -alkyl, C_3 - C_6 -cycloalkyl, C_2 - C_{10} -alkenyl, C_2 - C_{10} -alkinyl, C_1 - C_{10} -alkylcarbonyl, C_2 - C_{10} -alkenylcarbonyl, C_3 - C_{10} -alkinylcarbonyl, C_1 - C_{10} -alkylsulfonyl or C(R')=NOR", where the hydrocarbon radicals of these groups are unsubstituted or substituted by one to three radicals R^c :

R^c is cyano, nitro, amino, aminocarbonyl, aminothiocarbonyl, halogen, C₁-C₆-alkyl, C₁-C₆-haloalkyl, C₁-C₆-alkylsulfonyl, C₁-C₆-alkylsulfonyl, C₁-C₆-alkylsulfoxyl, C₁-C₆-alkoxy, C₁-C₆-haloalkoxy, C₁-C₆-alkylsulfoxyl, C₁-C₆-alkylthio, C₁-C₆-alkylamino, di-C₁-C₆-alkylamino, C₁-C₆-alkylaminocarbonyl, di-C₁-C₆-alkylaminocarbonyl, di-C₁-C₆-alkylaminothiocarbonyl, C₁-C₆-alkenyl, C₂-C₆-alkenyloxy, C₃-C₆-cycloalkyl, C₃-C₆-cycloalkyloxy, 5- or 6-membered heterocyclyl, 5- or 6-membered heterocyclyloxy, benzyl, benzyloxy, phenyl, phenoxy, phenylthio, 5- or 6-membered hetaryl, 5- or 6-membered hetaryloxy or hetarylthio, where the cyclic groups for their part may be partially of fully halogenated or may carry one to three radicals R^a; and

 R^3 is hydrogen, C_1 - C_6 -alkyl, C_2 - C_6 -alkenyl or C_2 - C_6 -alkinyl, where the hydrocarbon radicals of these groups may be unsubstituted or substituted by one to three radicals R^c ;

and

b) one or more ethylene modulators (II) selected from the group consisting of: oethylene biosynthesis inhibitors which inhibit the conversion of S-adenosyl-L-methionine into 1-aminocyclopropane-1-carboxylic acid (ACC), selected from derivatives of vinylglycine and hydroxylamines; oethylene biosynthesis inhibitors which block the conversion of ACC into ethylene, selected from the group consisting of: Co⁺⁺ or Ni⁺⁺ ions in plant-available forms; phenolic radical scavengers such as *n*-propyl gallate; polyamines, such as putrescine, spermine or spermidine; structural analogs of ACC, such as a-aminoisobutyric acid or Laminocyclopropene-1-carboxylic acid; salicylic acid or acibenzolar-Smethyl; structural analogs of ascorbic acid which act as inhibitors of ACC oxidase, such as prohexadione-Ca or trinexapac-ethyl; and triazolyl compounds such as paclobutrazol or uniconazole as inhibitors of cytochrome P-450-dependent monooxygenases whose main action is to block the biosynthesis of gibberellins;

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oinhibitors of the action of ethylene selected from the group consisting of: structural analogs of ethylene such as 1-methylcyclopropene or 2,5norbornadiene and 3-amino-1,2,4-triazole or Ag⁺⁺ ions

in a weight ratio of I to II of from 20:1 to 0.05:1.

- 2. (Original) A mixture as claimed in claim 1 where the compound of the formula I is a strobilurin derivative selected from the group consisting of azoxystrobin, dimoxystrobin, fluoxastrobin, kresoxim-methyl, metominostrobin, orysastrobin, trifloxystrobin, picoxystrobin or pyraclostrobin.
- 3. (Original) A mixture as claimed in claim 1 where the compound of the formula I is pyraclo-strobin.
- 4. (Original) A mixture as claimed in claim 1 where the ethylene modulators are Co^{++} ions, aminoethoxyvinylglycine, aminooxyacetic acid, prohexadione-Ca, trinexapacethyl, α -aminoisobutyric acid, salicylic acid or 3-amino-1,2,4-triazole.
- 5. (Original) A mixture as claimed in claim 1 where the ethylene modulators are Co⁺⁺ ions.

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6. (Original) A mixture as claimed in claim 1 where the ethylene modulators is prohexadione-Ca.

- 7. (Original) A mixture as claimed in claim 1 where the ethylene modulator is salicylic acid.
- 8. (Original) A mixture as claimed in claim 1 where the ethylene modulators are prohexadione-Ca together with Co⁺⁺ ions.
- 9. (Currently amended) A mixture as claimed in any of claims 1 to 8 claim 1 which additionally comprises an azole III selected from the group consisting of bromoconazole, cyproconazole, epoxiconazole, fenbuconazole, fluquiconazole, flugilazole, metconazole, myclobutanil, propiconazole, prochloraz, prothioconazole, tebuconazole or triticonazole.
- 10. (Currently amended) A mixture as claimed in any of claims 1 to 9 claim 1 which additionally comprises a surfactant selected from the group consisting of: polyoxyethylene sorbitan monolaurate, alkylphenoxy polyethoxy ethanol, fatty alcohol, fatty alcohol alkoxylate and sodium dodecylsulfate.

11. (Currently amended) A method for controlling rust infections in legumes, which comprises treating the above-ground plant parts of the legumes with an aqueous preparation of a mixture as claimed in any of claims 1 to 10 claim 1.

- 12. (Original) A process as claimed in claim 11, wherein rust infection on leaves and fruits of soya plants is controlled.
- 13. (Original) A process as claimed in claim 11, wherein the rust infection is caused by *Phakopsora* pachyrhizi and/or *Phakopsora meibomiae*.
- 14. (Currently amended) A process for increasing the yield and quality of legumes by using mixtures as claimed in any of claims 1 to 10 claim 1.
- 15. (Currently amended) A method for increasing the yield and quality of legumes applying an effective amount of a mixture as claimed in any of claims 1 to 10 claim 1.
- 16. (Currently amended) A method for reducing the ethylene evolution of plants by applying an effective amount of a mixture as claimed in claims 1 to 10 claim 1.
- 17. (Currently amended) A method for reducing undesired defoliation of crop plants by applying an effective amount of a mixture as claimed in claims 1 to 10 claim 1.

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18. (Original) A method for controlling harmful plant pathogens by applying an effective amount of Co⁺⁺ ions in plant-available form.